

Simona Cappelletti

List of Publications by Year in descending order

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Version: 2024-02-01

24
papers

572
citations

567281

15
h-index

642732

23
g-index

24
all docs

24
docs citations

24
times ranked

905
citing authors

#	ARTICLE	IF	CITATIONS
1	Cognitive development in Dravet syndrome: A retrospective, multicenter study of 26 patients. <i>Epilepsia</i> , 2011, 52, 386-392.	5.1	99
2	Panayiotopoulos syndrome: A clinical, EEG, and neuropsychological study of 93 consecutive patients. <i>Epilepsia</i> , 2010, 51, 2098-2107.	5.1	62
3	Defining the electroclinical phenotype and outcome of PCDH19-related epilepsy: A multicenter study. <i>Epilepsia</i> , 2018, 59, 2260-2271.	5.1	39
4	Electroencephalographic Features in Dravet Syndrome. <i>Journal of Child Neurology</i> , 2012, 27, 439-444.	1.4	34
5	Myoclonic astatic epilepsy: An age-dependent epileptic syndrome with favorable seizure outcome but variable cognitive evolution. <i>Epilepsy Research</i> , 2011, 97, 133-141.	1.6	33
6	Cognitive development in females with PCDH19 gene-related epilepsy. <i>Epilepsy and Behavior</i> , 2015, 42, 36-40.	1.7	32
7	CHD2 mutations are a rare cause of generalized epilepsy with myoclonic-atonic seizures. <i>Epilepsy and Behavior</i> , 2015, 51, 53-56.	1.7	28
8	PCDH19-related epilepsy and Dravet Syndrome: Face-off between two early-onset epilepsies with fever sensitivity. <i>Epilepsy Research</i> , 2016, 125, 32-36.	1.6	28
9	Symptoms of anxiety and depression and family's quality of life in children and adolescents with epilepsy. <i>Epilepsy and Behavior</i> , 2018, 79, 146-153.	1.7	28
10	PRRT2 is mutated in familial and non-familial benign infantile seizures. <i>European Journal of Paediatric Neurology</i> , 2013, 17, 77-81.	1.6	22
11	Cognitive Findings and Behavior in Children and Adolescents with Phenylketonuria. <i>Journal of Developmental and Behavioral Pediatrics</i> , 2013, 34, 392-398.	1.1	22
12	Mutation of <i>CHRNA2</i> in a family with benign familial infantile seizures: Potential role of nicotinic acetylcholine receptor in various phenotypes of epilepsy. <i>Epilepsia</i> , 2015, 56, e53-7.	5.1	19
13	Purified Cannabidiol for Treatment of Refractory Epilepsies in Pediatric Patients with Developmental and Epileptic Encephalopathy. <i>Paediatric Drugs</i> , 2019, 21, 283-290.	3.1	19
14	Epilepsy in ring 14 chromosome syndrome. <i>Epilepsy and Behavior</i> , 2012, 25, 585-592.	1.7	17
15	Epilepsy in Patients With Duplications of Chromosome 14 Harboring FOXG1. <i>Pediatric Neurology</i> , 2014, 50, 530-535.	2.1	17
16	Prolonged episode of dystonia and dyskinesia resembling status epilepticus following acute intrathecal baclofen withdrawal. <i>Epilepsy and Behavior</i> , 2011, 21, 321-323.	1.7	15
17	Ring 21 chromosome presenting with epilepsy and intellectual disability: Clinical report and review of the literature. , 2011, 155, 911-914.		13
18	Behavioral disorders as unusual presentation of pediatric extraventricular neurocytoma: report on two cases and review of the literature. <i>BMC Neurology</i> , 2014, 14, 242.	1.8	10

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19	Occipital seizures induced by Intermittent Photic Stimulation in Dravet syndrome. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2014, 23, 309-313.	2.0	10
20	Neonatal hemifacial spasm and fourth ventricle mass. <i>Developmental Medicine and Child Neurology</i> , 2012, 54, 697-703.	2.1	9
21	Acute intralesional recording in hypothalamic hamartoma: description of 4 cases. <i>Acta Neurologica Belgica</i> , 2015, 115, 233-239.	1.1	6
22	Comorbidities in Dravet Syndrome and Lennoxâ€Gastaut Syndrome. <i>SN Comprehensive Clinical Medicine</i> , 2021, 3, 2167-2179.	0.6	6
23	Ictal yawning in a patient with drug-resistant focal epilepsy: Video/EEG documentation and review of literature reports. <i>Epilepsy and Behavior</i> , 2011, 22, 602-605.	1.7	3
24	Structural Focal Temporal Lobe Seizures in a Child With Lipoproteinosis. <i>Pediatric Neurology</i> , 2015, 52, 104-106.	2.1	1